PHARMACY BOARD[657]

Notice of Intended Action

Twenty-five interested persons, a governmental subdivision, an agency or association of 25 or more persons may demand an oral presentation hereon as provided in Iowa Code section 17A.4(1)"b."

Notice is also given to the public that the Administrative Rules Review Committee may, on its own motion or on written request by any individual or group, review this proposed action under section 17A.8(6) at a regular or special meeting where the public or interested persons may be heard.

Pursuant to the authority of Iowa Code sections 124.301 and 147.76, the Board of Pharmacy hereby gives Notice of Intended Action to amend Chapter 22, "Unit Dose, Alternative Packaging, and Emergency Boxes," and Chapter 23, "Long-Term Care Pharmacy Practice," Iowa Administrative Code.

The amendments were approved at the August 29, 2012, regular meeting of the Board of Pharmacy.

The proposed amendments authorize a pharmacy other than a facility's primary provider pharmacy to provide to the facility to meet the needs of the facility's patients an emergency/first dose drug supply containing those drugs and products not stocked or available from the primary provider pharmacy. This additional supply may include, but is not limited to, parenteral or compounded drug products.

Requests for waiver or variance of the discretionary provisions of Board rules will be considered pursuant to 657—Chapter 34.

Any interested person may present written comments, data, views, and arguments on the proposed amendments not later than 4:30 p.m. on October 23, 2012. Such written materials may be sent to Terry Witkowski, Executive Officer, Board of Pharmacy, 400 S.W. Eighth Street, Suite E, Des Moines, Iowa 50309-4688; or by e-mail to terry.witkowski@iowa.gov.

After analysis and review of this rule making, no impact on jobs has been found.

These amendments are intended to implement Iowa Code sections 124.301, 155A.13, and 155A.15. The following amendments are proposed.

ITEM 1. Amend subrule 22.7(1) as follows:

22.7(1) Emergency/first dose drug supplies. All contents Contents of the emergency/first dose drug supply shall be provided by one a primary provider pharmacy designated by the facility, and the drug supply shall be available to meet the needs of all patients of the facility, without penalty or discrimination. A second provider pharmacy may provide an emergency/first dose drug supply consisting only of drugs and products not stocked or available from the primary provider pharmacy including, but not limited to, parenteral or compounded drug products. The provider pharmacy pharmacies shall be properly registered with the federal Drug Enforcement Administration (DEA) and the board and shall be currently licensed by the board. The provider pharmacist or pharmacists, the consultant pharmacist, the director of nursing of the facility, and the medical director of the facility, or their respective designees, shall jointly determine and prepare a list of drugs necessary for prompt use in patient care that will be available in the each emergency/first dose drug supply. Drugs shall be listed by identity and quantity, shall be limited to drugs necessary to meet the emergency needs of the patients served, and shall be periodically reviewed pursuant to policy. Careful patient planning should be a cooperative effort between the pharmacy pharmacies and the facility to make drugs available, and this supply emergency/first dose drug supplies shall only be used for emergency or unanticipated needs. The intent of the emergency/first dose drug supply is not to relieve a pharmacy of the responsibility for timely provision of a patient's routine drug needs; the intent is to ensure that a supply of drugs is available to each patient in case of urgent need. The drugs in the emergency/first dose drug supply supplies are the responsibility of the respective provider pharmacy and, therefore, shall not be used or altered in any way except as provided in this rule.

ITEM 2. Amend rule 657—23.5(124,155A) as follows:

657—23.5(124,155A) Emergency drugs. A supply of emergency drugs may be provided by one <u>or more</u> long-term care pharmacy provider pharmacies to the facility pursuant to rule 657—22.7(124,155A).

- **23.5(1)** Emergency medication order—pharmacist review. When an emergency drug is provided pursuant to rule 657—22.7(124,155A), the medication order shall be reviewed by the resident's dispensing pharmacist prior to the administration of a second dose.
- 23.5(2) Other emergency drugs and devices. In addition to an one or more emergency box boxes or stat drug box boxes, a long-term care facility staffed by one or more persons licensed to administer drugs may maintain a stock of intravenous fluids, irrigation fluids, heparin flush kits, medicinal gases, sterile water and saline, and prescription devices. Such stock shall be limited to a listing to be determined by the provider pharmacist in consultation with the consultant pharmacist and the medical director and director of nursing of the facility.